In the claims:

- 1. (currently amended) A method of treating a stenosis or restenosis in a coronary blood vessel <u>characterized by a proximal end nearest the origination of blood flow from the heart and a distal end, said method comprising the steps of:</u>
 - implanting a stent at a first location within the coronary blood vessel; and
 - injecting a therapeutic agent comprising an anti-restenosis agent into the myocardium proximate the coronary blood vessel at a second location, where the second location is distal, in relation to the coronary blood vessel, to the first location at a site distal to the stent.
- 2. (original) The method of claim 1 further comprising the steps of:
 - injecting the therapeutic agent into the myocardium from an endocardial space of the heart.
- 3. (original) The method of claim 1 further comprising the steps of:
 - injecting the therapeutic agent peri-adventitially through the blood vessel wall.
- 4. (original) The method of claim 1 further comprising the step of:
 - injecting the therapeutic agent peri-adventitially through a coronary vein or coronary sinus.
- 5. (canceled)

- 6. (original) The method of claim 1, 2, 3 or 4 further comprising the steps of:
 - selecting the anti-restenosis agent from the group comprising anti-oxidant drugs, anti-inflammatory drugs, anti-neoplastic agents, anti-angiogenic agents and gene therapy agents.
- 7. (original) The method of claim 1, 2, 3 or 4 further comprising the step of:
 - providing the therapeutic agent in a time release formulation.
- 8. (original) The method of claim 1, 2, 3 or 4 further comprising the step of:
 - providing the therapeutic agent in a microsphere formulation.
- 9. (original) The method of claim 1, 2, 3 or 4 further comprising the step of:
 - providing the therapeutic agent in a formulation in which the therapeutic agent is encapsulated in micelles.
- 10. (original) The method of claim 1, 2, 3 or 4 further comprising the step of:
 - providing the therapeutic agent in a formulation in which the therapeutic agent is encapsulated in liposomes.
- 11. (currently amended) A method of treating a stenosis or restenosis in a coronary blood vessel characterized by a

proximal end nearest the origination of blood flow from the heart and a distal end, said method comprising the steps of:

performing an angioplasty procedure <u>at a first location</u>
within the coronary blood vessel; and

- injecting a therapeutic agent comprising an anti-restenosis agent into the myocardium proximate the coronary blood vessel at a second location , where the second location is distal, in relation to the coronary blood vessel, to the first location at a site distal to the site of angioplasty.
- 12. (original) The method of claim 11 further comprising the steps of:
 - injecting the therapeutic agent into the myocardium from an endocardial space of the heart.
- 13. (original) The method of claim 11 further comprising the steps of:
 - injecting the therapeutic agent peri-adventitially through the blood vessel wall.
- 14. (original) The method of claim 11 further comprising the step of:
 - injecting the therapeutic agent peri-adventitially through a coronary vein or coronary sinus.
- 15. (canceled)
- 16. (original) The method of claim 11, 12, 13 or 14 further comprising the steps of:

- selecting the anti-restenosis agent from the group comprising anti-oxidant drugs, anti-inflammatory drugs, anti-neoplastic agents, anti-angiogenic agents and gene therapy agents.
- 17. (original) The method of claim 11, 12, 13 or 14 further comprising the step of:

providing the therapeutic agent in a time release formulation.

18. (original) The method of claim 11, 12, 13 or 14 further comprising the step of:

providing the therapeutic agent in a microsphere formulation.

19. (original) The method of claim 11, 12, 13 or 14 further comprising the step of:

providing the therapeutic agent in a formulation in which the therapeutic agent is encapsulated in micelles.

20. (original) The method of claim 11, 12, 13 or 14 further comprising the step of:

providing the therapeutic agent in a formulation in which the therapeutic agent is encapsulated in liposomes.

21. (currently amended) A method of treating a segment of a coronary blood vessel, the segment characterized by a proximal end nearest the origination of blood flow from the heart and a distal end, said method comprising the steps of:

- injecting a therapeutic agent comprising an anti-restenosis agent into the myocardium proximate the coronary blood vessel at a site distal, to the segment to be treated.
- 22. (original) The method of claim 21 further comprising the steps of:
 - injecting the therapeutic agent into the myocardium from an endocardial space of the heart.
- 23. (original) The method of claim 21 further comprising the steps of:
 - injecting the therapeutic agent peri-adventitially through the blood vessel wall.
- 24. (original) The method of claim 21 further comprising the step of:
 - injecting the therapeutic agent peri-adventitially through a coronary vein or coronary sinus.
- 25. (canceled)
- 26. (original) The method of claim 21, 22, 23 or 24 further comprising the steps of:
 - selecting the anti-restenosis agent from the group comprising anti-oxidant drugs, anti-inflammatory drugs, anti-neoplastic agents, anti-angiogenic agents and gene therapy agents.
- 27. (original) The method of claim 21, 22, 23 or 24 further comprising the step of:

- providing the therapeutic agent in a time release formulation.
- 28. (original) The method of claim 21, 22, 23 or 24 further comprising the step of:
 - providing the therapeutic agent in a microsphere formulation.
- 29. (original) The method of claim 21, 22, 23 or 24 further comprising the step of:
 - providing the therapeutic agent in a formulation in which the therapeutic agent is encapsulated in micelles.
- 30. (original) The method of claim 21, 22, 23 or 24 further comprising the step of:
 - providing the therapeutic agent in a formulation in which the therapeutic agent is encapsulated in liposomes.
- 31. (currently amended) A method of treating \underline{a} segment of a coronary blood vessel characterized by an intraluminal disease, the segment further characterized by a proximal end nearest the origination of blood flow from the heart and a distal end, said \underline{method} comprising the steps of:
 - injecting a therapeutic agent into the myocardium proximate the coronary blood vessel at a site distal, to the segment to be treated.
- 32. (original) The method of claim 31 further comprising the steps of:

- injecting the therapeutic agent into the myocardium from an endocardial space of the heart.
- 33. (original) The method of claim 31 further comprising the steps of:
 - injecting the therapeutic agent peri-adventitially through the blood vessel wall.
- 34. (original) The method of claim 31 further comprising the step of:
 - injecting the therapeutic agent peri-adventitially through a coronary vein or coronary sinus.
- 35. (canceled)
- 36. (original) The method of claim 31, 32, 33 or 34 further comprising the steps of:
 - selecting the therapeutic agent from the group comprising anti-oxidant drugs, anti-inflammatory drugs, antineoplastic agents, anti-angiogenic agents and gene therapy agents.
- 37. (original) The method of claim 31, 32, 33 or 34 further comprising the step of:
 - providing the therapeutic agent in a time release formulation.
- 38. (original) The method of claim 31, 32, 33 or 34 further comprising the step of:

providing the therapeutic agent in a microsphere formulation.

39. (original) The method of claim 31, 32, 33 or 34 further comprising the step of:

providing the therapeutic agent in a formulation in which the therapeutic agent is encapsulated in micelles.

40. (original) The method of claim 31, 32, 33 or 34 further comprising the step of:

providing the therapeutic agent in a formulation in which the therapeutic agent is encapsulated in liposomes.

- 41. (currently amended) A kit for delivering a therapeutic agent to a patient suffering from vascular disease characterized by a diseased treatment region in a coronary blood vessel, the diseased treatment region characterized by a proximal end nearest the origination of blood flow from the heart and a distal end, said kit comprising:
 - a catheter having means for introducing a therapeutic agent into in a perivascular space surrounding the blood vessel; and
 - a dose of therapeutic agent suitable for introduction into the perivascular space surrounding the blood vessel through the catheter;
 - instructions for use of the catheter according to the following method:

positioning the means for introducing into the perivascular space; and

delivering a dose of the therapeutic agent into the perivascular space near the diseased treatment region at a site distal, to the diseased treatment region.